

Clstn3 Cas9-KO Strategy

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Project Overview



Project Name

Clstn3

Project type

Cas9-KO

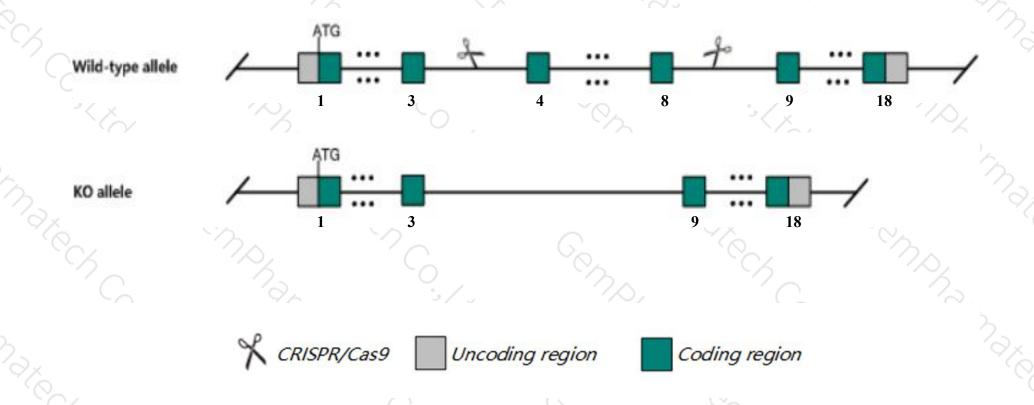
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Clstn3 gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Clstn3* gene has 9 transcripts. According to the structure of *Clstn3* gene, exon4-exon8 of *Clstn3-201*(ENSMUST00000008297.4) transcript is recommended as the knockout region. The region contains 940bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Clstn3* gene. The brief process is as follows: CRISPR/Cas9 system

Notice



- > According to the existing MGI data, mice homozygous for a knock-out allele exhibit reductions in excitatory and inhibitory synapse density and deficits in synaptic transmission.
- ➤ Transcript *Clstn3-203, Clstn3-206 and Clstn3-208* may not be affected.
- > The *Clstn3* gene is located on the Chr6. If the knockout mice are crossed with other mice strains to obtain double gene positive homozygous mouse offspring, please avoid the two genes on the same chromosome.
- This strategy is designed based on genetic information in existing databases. Due to the complexity of biological processes, all risk of the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Gene information (NCBI)



Clstn3 calsyntenin 3 [Mus musculus (house mouse)]

Gene ID: 232370, updated on 13-Mar-2020

Summary

☆ ?

Official Symbol Clstn3 provided by MGI

Official Full Name calsyntenin 3 provided by MGI

Primary source MGI:MGI:2178323

See related Ensembl: ENSMUSG00000008153

Gene type protein coding
RefSeq status VALIDATED
Organism Mus musculus

Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;

Muroidea; Muridae; Murinae; Mus; Mus

Also known as CSTN3, Clstn3b, Cs3, Cst-3, alc-beta

Expression Biased expression in cerebellum adult (RPKM 97.0), cortex adult (RPKM 62.3) and 8 other tissuesSee more

Orthologs <u>human</u> all

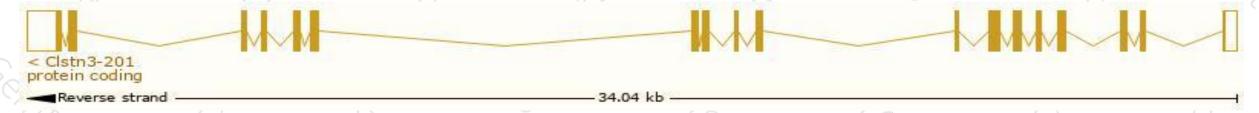
Transcript information (Ensembl)



The gene has 9 transcripts, all transcripts are shown below:

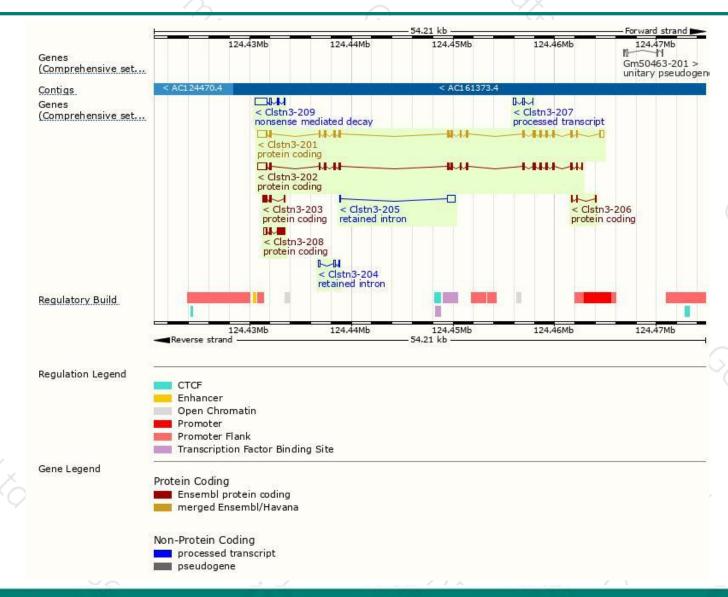
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
CIstn3-201	ENSMUST00000008297.4	4014	956aa	Protein coding	CCDS20519	Q99JH7	TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P3
Cistn3-202	ENSMUST00000112523.7	3703	919aa	Protein coding	CCDS85156	D3Z601	TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS ALT2
CIstn3-208	ENSMUST00000238807.1	1382	357aa	Protein coding	-	A0A4P8DYG6	GENCODE basic
CIstn3-203	ENSMUST00000124998.1	728	<u>191aa</u>	Protein coding	-	127	TSL:2 GENCODE basic
CIstn3-206	ENSMUST00000150774.2	366	64aa	Protein coding	-	A0A0N4SW90	CDS 3' incomplete TSL:5
Clstn3-209	ENSMUST00000239139.1	1687	78aa	Nonsense mediated decay	8 .	-	
CIstn3-207	ENSMUST00000156040.1	391	No protein	Processed transcript	-	(39)	TSL:2
CIstn3-205	ENSMUST00000147947.1	765	No protein	Retained intron	20		TSL:2
CIstn3-204	ENSMUST00000143283.1	474	No protein	Retained intron	-		TSL:3

The strategy is based on the design of *Clstn3-201* transcript, the transcription is shown below:



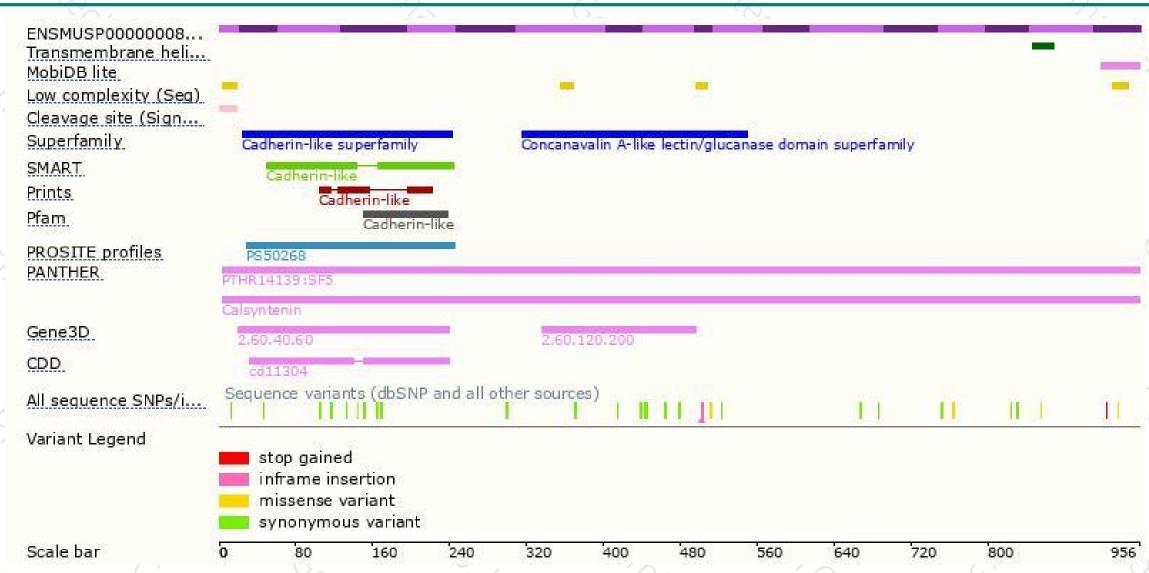
Genomic location distribution





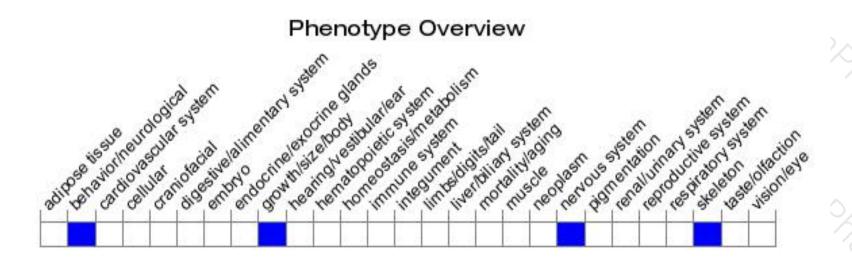
Protein domain





Mouse phenotype description(MGI)





Phenotypes affected by the gene are marked in blue.Data quoted from MGI database(http://www.informatics.jax.org/).

According to the existing MGI data, mice homozygous for a knock-out allele exhibit reductions in excitatory and inhibitory synapse density and deficits in synaptic transmission.



If you have any questions, you are welcome to inquire. Tel: 400-9660890





